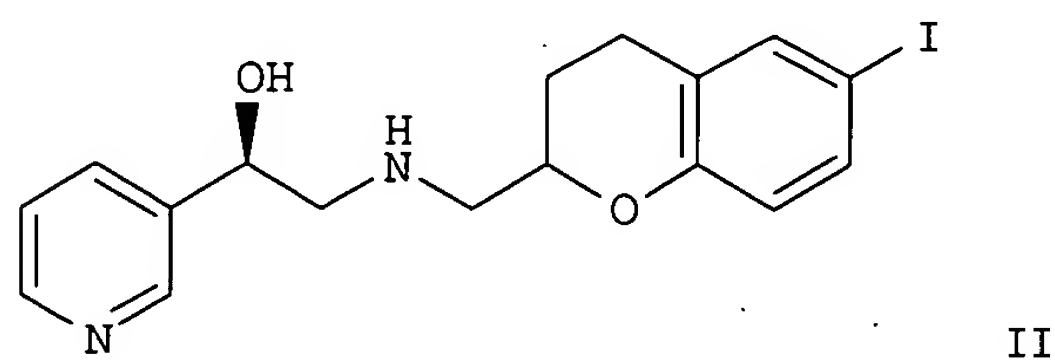
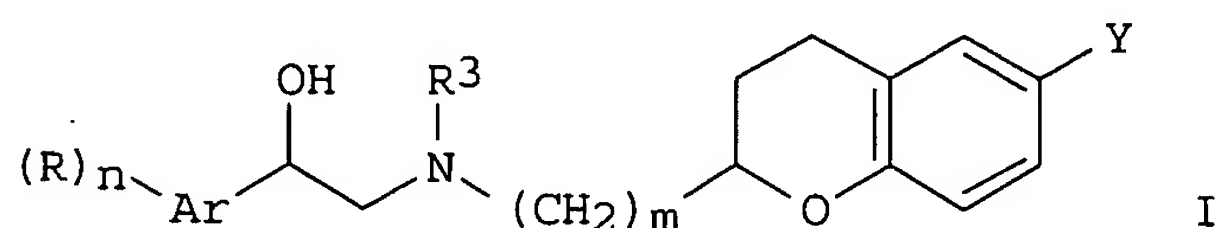


AN 1996:646308 CAPLUS  
 DN 125:300822  
 TI Preparation of N-**chromanyl** and N-**chromanylmethyl**  
 2-amino-1-phenylethanol compounds as adrenergic  $\beta$ 3-receptor  
 stimulants  
 IN Tsucha, Susumu; Fukuzaki, Atsushi; Takenawa, Noriko; Ozaka, Kazuya  
 PA Tokyo Tanabe Co, Japan  
 SO Jpn. Kokai Tokkyo Koho, 10 pp.  
 CODEN: JKXXAF  
 DT Patent  
 LA Japanese  
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| PRAI | JP 1995-6912      |      | 19950120 |                 |          |
| OS   | MARPAT 125:300822 |      |          |                 |          |



AB This invention relates to novel 2,6-substituted **chroman** derivs. which are useful in the treatment of **.beta.3-adrenoreceptor** mediated conditions. Title compds. I [wherein R = independently OH, :O, halo, CN, NO<sub>2</sub>, (halo)alkyl, CF<sub>3</sub>, NR<sub>1</sub>R<sub>1</sub>, SR<sub>1</sub>, OR<sub>1</sub>, SO<sub>2</sub>R<sub>2</sub>, OCOR<sub>2</sub>, NR<sub>1</sub>COR<sub>2</sub>, COR<sub>2</sub>, NR<sub>1</sub>SO<sub>2</sub>R<sub>2</sub>, or (un)substituted Ph or heterocyclyl; R<sub>1</sub> = independently H, (CH<sub>2</sub>)<sub>m</sub>O(CH<sub>2</sub>)<sub>m</sub>R<sub>5</sub>, or (un)substituted (cyclo)alkyl, Ph, or naphthyl; or NR<sub>1</sub>R<sub>1</sub> = heterocyclyl; R<sub>2</sub> = independently R<sub>1</sub>, OR<sub>1</sub>, NR<sub>1</sub>R<sub>1</sub>, or (un)substituted NHSO<sub>0</sub>-2-Ph, NHSO<sub>0</sub>-2-naphthyl, NHSO<sub>0</sub>-2-alkyl, or heterocyclyl; R<sub>3</sub> = H, alkyl, or COR<sub>3</sub>; R<sub>4</sub> = H, alkyl(phenyl), or alkylpyridyl; R<sub>5</sub> = H or CO<sub>2</sub>H; R<sub>6</sub> = H or (un)substituted alkyl or alkyl-SO<sub>0</sub>-2-alkyl; Ar = Ph or (fused) hetero(aryl); Y = halo, NO<sub>2</sub>, R<sub>6</sub>, SR<sub>1</sub>, SO<sub>0</sub>-2C<sub>6</sub>H<sub>4</sub>CO<sub>2</sub>R<sub>1</sub>, (CONR<sub>4</sub>CR<sub>4</sub>R<sub>4</sub>)pCO<sub>2</sub>R<sub>1</sub>, or (un)substituted Ph or heterocyclyl; m = 1-3; n = 0-5; p = 1 or 2; and pharmaceutically acceptable salts and esters thereof] were prepared as  $\beta$  3-**adrenoceptor** agonists. For example, coupling of (2R)-6-iodo-3,4-dihydro-2H-chromene-2-carboxylic acid and (1R)-2-amino-1-(3-pyridinyl)ethanol•2HCl with 1-hydroxybenzotriazole, 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide•HCl, and TEA in CH<sub>2</sub>Cl<sub>2</sub> gave the amide (74%). Reduction using borane-dimethylsulfide complex in THF afforded the **chromanmethaneamine** II (84%). Over one hundred compds. of the invention demonstrated **.beta.3-adrenergic** receptor agonist activity with EC<sub>50</sub> values  $\leq$  1 $\mu$ M. I are useful in the treatment of **.beta.3-adrenergic** receptor mediated conditions, including obesity, diabetes, **gastrointestinal** disorders, cardiovascular disorders, and urinary disorders (no data).